CLAIMS

What is claimed is:

1. A hERG channel-expressing cell population comprising cells capable of expressing a channel of which the hERG current as determined by patch clamping with a fully automated high throughput patch clamp system is 0.6 nA or more, wherein the proportion ratio of said cells is 40% or more relative to the total number of hERG gene-transferred cells within said population.—

2. The cell population according to claim 1, wherein the hERG gene has been transferred with a virus vector.

- 3. The cell population according to claim 2, wherein the virus vector is a retrovirus vector or a lentivirus vector.
 - 4. The cell population according to any one of claims 1 to 3 claim 1, wherein the average value of the hERG current in the total cell population cells is 0.3 nA or more.
- 5. A cell capable of expressing a hERG channel of which the hERG current as determined by patch clamping with a fully automated high throughput patch clamp system is 1.0 nA or more.—
- 6. The cell according to claim 5, wherein the hERG gene has been transferred with a virus vector.
 - 7. The cell according to claim 6, wherein the virus vector is a retrovirus vector or a lentivirus vector.
- 8. A method of preparing the cell population according to any one of claims 1 to 4 or the cell according to any one of claims 5 to 7claim 1, the method comprising which comprises expressing hERG channels using via a virus vector.
- The method according to claim 8, wherein the virus vector is a retrovirus vector or a
 lentivirus vector.

- 10. (Canceled)The method according to claim 8, wherein the virus-vector is a retrovirus vector:
- 5 11. The method according to <u>claim 8 any one of claims 8 to 10</u>, the method further <u>comprising thewhich comprises a</u> step of concentrating the virus vector by ultracentrifugation.
- 12. A method of measuring hERG current inhibitory activity, the method comprising which eemprises using the cell population according to any one of claims 1 to 4 or the cell according to any one of claims 5 to 7 claim 1.—

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- 13. The method according to claim 12, the method further comprisingwhich comprises using a fully automated high throughput patch clamp system.
- 14. A method of measuring hERG current inhibitory activity, the method comprising which comprises using a cell population or a cell prepared by the method according to any one of claims 8.9, orto 11 claim 8.
- 20 15. The method according to claim 14, the method further comprising which comprises using a fully automated high throughput patch clamp system.
 - 16. A method of screening a compoundfor compounds, or a saltsalts thereof for its hERG current altering effect, that alter or not alter hERG currents, the method comprising which comprises using the cell population according to any one of claims 1 to 4 or the cell according to any one of claims 5 to 7claim 1.—
 - 17. The method according to claim 16, the method further comprising which comprises using a fully automated high throughput patch clamp system.
 - 18. A method of screening a compound for compounds, or a saltsalts thereof for its hERG current altering effect, that alter or not alter hERG currents, the method comprising which comprises using a cell population or a cell prepared by the method according to any one of claims 8 9, or to 11 claim 8.

- 19. The method according to claim 18, the method further comprising which comprises using a fully automated high throughput patch clamp system.—
- 20. A method of measuring hERG current inhibitory activity, the method comprising using
 the cell population according to claim 5.
 - 21. A method of screening a compound or a salt thereof for its hERG current altering effect, the method comprising using the cell population according to claim 5.